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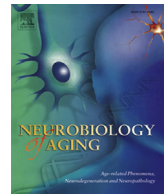
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## Functional connectivity among brain regions affected in Alzheimer's disease is associated with CSF TNF- $\alpha$ in APOE4 carriers



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### ABSTRACT

It is now recognized that understanding how neuroinflammation affects brain function may provide new insights into Alzheimer's pathophysiology. Tumor necrosis factor (TNF)- $\alpha$ , an inflammatory cytokine marker, has been implicated in Alzheimer's disease (AD), as it can impair neuronal function through suppression of long-term potentiation. Our study investigated the relationship between cerebrospinal fluid TNF- $\alpha$  and functional connectivity (FC) in a cohort of 64 older adults ( $\mu$  age = 69.76 years; 30 cognitively normal, 34 mild AD). Higher cerebrospinal fluid TNF- $\alpha$  levels were associated with lower FC among brain regions important for high-level decision-making, inhibitory control, and memory. This effect was moderated by apolipoprotein E- $\epsilon$ 4 (APOE4) status. Graph theory metrics revealed there were significant differences between APOE4 carriers at the node level, and by diagnosis at the network level suggesting global brain network dysfunction in participants with AD. These findings suggest proinflammatory mechanisms may contribute to reduced FC in regions important for high-level cognition. Future studies are needed to understand the role of inflammation on brain function and clinical progression, especially in APOE4 carriers.

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### 1. Introduction

Beta-amyloid and neurofibrillary tangles are the hallmark features of Alzheimer's pathology. However, it is increasingly recognized that other pathological insults, such as neuroinflammation, may be involved in the Alzheimer's cascade (Ardura-Fabregat et al., 2017; Bronzuoli et al., 2016; Sweeney et al., 2015). For instance, activation of neuroinflammatory systems is elevated in Alzheimer's disease (AD) relative to normal aging (Breunig et al., 2013; Heneka et al., 2015). Microglia, astrocytes, and even neurons have been shown to directly react and contribute to chronic neuroinflammatory

responses in AD. It has been suggested that astrocyte and microglial activation may be an early event in the pathophysiology of AD, occurring even in the absence of focal amyloid- $\beta$  (A $\beta$ ) deposition (Dudal et al., 2004).

Apolipoprotein E- $\epsilon$ 4 (APOE4), a major genetic risk factor for late-onset AD, provides a unique opportunity to evaluate AD risk before the onset of clinical symptoms. APOE is an important modulator of immune responses, as evidenced by greater immune activation in mice lacking APOE (Liu et al., 2015). Particularly, APOE4 has been shown to promote the strongest proinflammatory effects in mouse models (Guo et al., 2004; Vitek et al., 2009) with additional evidence that APOE4 transgenic mice show increased susceptibility to inflammation compared with APOE2 and APOE3 mice (den Hartigh et al., 2012; Rodriguez et al., 2014; Zhu et al., 2012).

The induction of a microglial neuroinflammatory response results in the release of various inflammatory mediators including an array of neurotoxic cytokines (Akiyama et al., 2000; Tan et al., 2002).

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Notably, tumor necrosis factor- $\alpha$  (TNF- $\alpha$ ) has been shown to be markedly elevated in patients with AD (Alsadany et al., 2013; Heneka and O'Banion, 2007; Liu et al., 2014), suggesting a chronic neuro-inflammatory state that intensifies the secretion of associated cytokines and the duration of the immune response, which together may impact neuronal health. In fact, TNF- $\alpha$ , a member of the cytokine family, can directly impair neuronal function and suppress hippocampal long-term potentiation (LTP), a mechanism essential for memory storage and consolidation (Cunningham et al., 1996; Tancredi et al., 1992). For instance, young APPV7171 transgenic mice show significantly decreased hippocampal LTP preceding amyloid plaque deposition (Heneka et al., 2005).

In human studies, elevated glial activation is associated with increased inflammatory cytokine activity including TNF- $\alpha$  in patients with mild to moderate dementia compared with healthy individuals (Cagnin et al., 2001; Paganelli et al., 2002; Tarkowski et al., 2003). Furthermore, in a neuropsychiatric population, elevated inflammation, as measured by other cytokines such as TNF- $\alpha$  and C-reactive protein (CRP) in plasma, has been associated with decreased cortical-striatal connectivity among those diagnosed as clinically depressed compared with normal adults (Felger et al., 2016). These studies suggest that cytokines such as TNF- $\alpha$  may directly impair neuronal function, either by measurement of LTP suppression or decreased functional connectivity (FC) and that early focal inflammatory events may contribute to neuronal dysfunction well before cell death. Activated glial cells are believed to modulate synaptic transmission (Bacci et al., 1999; Eroglu and Barres, 2010), and conversely, synaptic activity can induce activation of neuroimmune cells, resulting in neurogenic neuroinflammation (Tian et al., 2012). Collectively, these studies demonstrate a relationship between neuroinflammation and neural function, with chronic neuroinflammation leading to the loss of synapse associated proteins, thereby causing neuronal damage (Ma et al., 2014).

Although this evidence implicates a role of neuroinflammation in AD pathogenesis, little is known about the effect of TNF- $\alpha$  on brain FC. Large-scale changes in global brain function are known to be associated with early changes in the AD cascade using resting-state functional magnetic resonance imaging (rsfMRI) and brain network connectomics (Damoiseaux et al., 2012; Supek et al., 2008). AD pathophysiology has long been associated with impaired synchrony among regions important for cognition. Recently, studies using rsfMRI in a brain connectomic framework have shown aberrant FC in the earliest stages of AD, concluding that abnormal FC may be one of the earliest indicators of neurodegeneration (Contreras et al., 2017, 2019). The aim of the present study was to first investigate a relationship between cerebrospinal fluid (CSF) inflammatory marker TNF- $\alpha$  and rsfMRI FC among our cohort, and, second, determine if APOE4 carrier status influences this relationship. Last, we hope to better characterize any differences based on APOE4 carrier status and clinical diagnosis using graph theory metrics.

## 2. Materials and methods

### 2.1. Participants

Our prospective study sample consisted of 64 older participants who were recruited through the Knight Alzheimer's Disease Research Center at Washington University at St. Louis. Study procedures were approved by the local Institutional Review Board and performed in accordance with the 1964 Declaration of Helsinki. Informed consent was obtained from all participants before study enrollment. Clinical diagnosis for each participant was based on standard criteria as incorporated in the Uniform Data Set (Morris et al., 2006). Participants were included based on availability of T1-weighted MPRAGE scan, rsfMRI scan, and CSF biomarker data.

Specifically, there were 91 subjects with available CSF markers. Of those 91 subjects, 64 had available rsfMRI data. Therefore, we selected all of the 64 subjects with both CSF markers and rsfMRI to be included in this study. All biomarker assays and quantitative MRI scans were conducted by investigators blinded to the clinical status of the participant.

### 2.2. CSF biomarkers and genotyping

#### 2.2.1. Inflammatory markers

Meso Scale Discovery (MSD) multiplex assay was used to determine CSF levels for TNF- $\alpha$  (Cat. No. K15049G, MSD, Rockville, MD, USA).

#### 2.2.2. Amyloid- $\beta$ peptide

Enzyme-linked immunosorbent assay was used to determine CSF A $\beta$ 42 in all participants (Cat. No. 81583, Innotech, Belgium).

#### 2.2.3. Tau

MSD assay was used to determine CSF levels of total tau on all participants (Cat. No. K151LAE, MSD, Rockville, MD, USA), and enzyme-linked immunosorbent assay was used to determine phosphorylated tau (pT181) (Cat. No. 81581, Innotech, Belgium).

#### 2.2.4. APOE genotyping

APOE genotyping was performed (as outlined in Nation et al. [2019]). Participants with at least one copy of the E4 gene were considered APOE4 carriers. There were no E2 homozygous or heterozygous carriers.

### 2.3. MRI data acquisition, processing, and analyses

#### 2.3.1. Brain imaging acquisition

All images were obtained on a Siemens TrioTim 3T scanner using a 20-channel head coil. Anatomical T1-weighted 3D gradient echo pulse sequence scans were acquired with the following parameters: flip angle = 8°, TR/TE = 2400/3.16 ms, FOV 256 × 256 mm, voxel size was 1.1 × 1.1 × 1.2 mm<sup>3</sup> isotropic, length of scan = 7.04 minutes. rsfMRI scans were acquired eyes closed with the following parameters: flip angle 90°, TR/TE = 2200/27 ms, FOV 384 × 384 mm, voxel size: 4 mm<sup>3</sup> isotropic, scan time = 6.01 minutes.

#### 2.3.2. Preprocessing and correlation analysis

rsfMRI images were preprocessed using the CONN-toolbox v18a ((Whitfield-Gabrieli and Nieto-Castanon, 2012): <http://www.nitrc.org/projects/conn>) in SPM12 for data analysis. The default preprocessing pipeline was utilized in which case, all functional images were realigned (motion corrected), centered, slice time—corrected, corrected for motion artifacts using the artifact detection tools, and coregistered to their corresponding structural images. Images were identified as outliers if the head movement (in direction x, y, z) was more than 5 standard deviations from the mean intensity of the entire run, or a 97th percentile threshold. All structural images were then centered and segmented into cerebrospinal fluid, gray and white matter, and spatially normalized to the Montreal Neurological Institute template. Functional images were then normalized to Montreal Neurological Institute space using the deformation field from the corresponding structural images and spatially smoothed to allow for better registration and reduction of noise using a 6 mm FWHM Gaussian kernel.

After preprocessing, the CompCor strategy (Behzadi et al., 2007) was implemented to account for white matter and CSF noise using principal component analysis. The analyses did not include global signal regression to avoid potential false anticorrelations (Murphy et al., 2009). Motion parameters, cerebrospinal fluid, and white

matter were included in the model and considered as variables of no interest. The mean BOLD signal time course was then extracted from every ROI and band pass filtered (0.009–0.09 Hz). To define our ROIs, we used the default-atlas implemented in the CONN toolbox (v18a) that uses a combination of the Harvard-Oxford atlas and the AAL atlas as well as a few commonly used ROIs from well-known networks and areas based of the work of Yeo et al. (Yeo et al., 2011) and excluded cerebellar and primary sensory areas. Pearson's correlation coefficients were calculated for all pairwise comparisons between ROIs (see [supplemental material, Appendix](#) for complete list of ROIs used).

### 2.3.3. *TNF- $\alpha$ -specific functional connectivity analysis and selection of seed region*

To explore all possible associations between TNF- $\alpha$  and ROI to ROI functional connections, linear regression was used to compute the correlation between all pairwise ROIs FC strength and TNF- $\alpha$ . This allowed us to look at the matrix of effects between using all ROIs as potential “seeds” and all selected target ROIs while also correcting for multiple comparisons, using connection-level threshold (e.g., analysis-level FDR  $p < 0.05$ ). Based on this analysis, the seed region was defined as the node with the strongest link between all ROIs and TNF- $\alpha$ , revealing the nucleus accumbens (NAcc). Other regions implicated in AD, such as the hippocampus and parahippocampus, were not found to be associated with TNF- $\alpha$  at FDR-corrected  $p$ -value  $< 0.05$ . In our analysis, we investigated regions functionally connected to either the L or R NAcc as seed region as well as investigating regions functionally connected to both L and R regions, which revealed similar results. Because there was no prior evidence to suggest left or right NAcc differed in their functional or anatomical role we chose to report on regions functionally connected to either the L or R NAcc as seed region. Age, sex, education, and diagnosis were included as model covariates, and all analyses were FDR-corrected ( $p < 0.05$ ) for multiple comparisons. Moderation analysis was carried out using the PROCESS macro (Hayes, 2017) version 3.1 in SPSS version 25 to assess if the association between TNF- $\alpha$  and FC was moderated by APOE4 status.

### 2.3.4. *Graph-based network analysis*

Graph theory metrics including average path length, clustering coefficient, local efficiency and global efficiency were calculated using CONN toolbox to characterize the dynamics of FC within our cohort. These measures were calculated at the node level for our seed regions (NAcc left and right) as well as at the whole-brain network level using all ROIs listed in [supplemental table 1](#). A graph adjacency matrix  $A(i, j)$  was computed by thresholding an ROI-to-ROI correlation matrix  $r(i, j)$  for each subject. To obtain a whole-brain network value for each metric for each subject, we averaged metric ROI values across the brain for each subject. Average path length was defined as the average of the shortest paths between each ROI and all other ROIs in the graph. Clustering coefficient, a measure of the degree to which nodes in a graph tend to cluster together, was calculated as the fraction of edges among all possible edges in the local neighboring subgraph for our seed ROI. Local efficiency represents a measure of local integration, characterizing the degree of interconnectedness of a node. Global efficiency, a measurement of node centrality within a network characterizing the global connectedness, was defined as the average of inverse distances between each ROI and all other ROIs in the graph. A detailed description of equations for graph theory metrics can be found at [www.nitrc.org/projects/conn](http://www.nitrc.org/projects/conn).

Partial correlation analysis was used to assess the relationship at the node level between graph theory metrics and TNF- $\alpha$  levels in APOE4 carriers and noncarriers and between diagnoses. One-way ANOVA was used to compare all metrics at the whole-brain

network level between both APOE4 carriers and noncarriers as well as between diagnoses.

Last, the variance, skewness, and kurtosis were calculated to help characterize the group differences between AD and cognitively normal (CN) participants for the aforementioned graph theory metrics. The variance is the expectation of the squared deviation of the observation from the sample mean and therefore is concerned with the “spread” of the sample. Skewness is concerned with symmetry of the sample. A skewness of 0 means complete symmetry, a negative skew indicates a left tail, and a positive skew indicates a longer right tail. Kurtosis was used to measure the sharpness of the sample peak with higher values indicating a more well-defined mean and more observations around the tail(s). Smaller values indicate flatter distributions.

## 3. Results

### 3.1. *Study participants*

64 participants (30 CN, 34 mild AD) were included in this study. Of the total sample, 59% were men and 55% were APOE4 carriers. Participants were well educated with an average of 15.19 years of education. APOE4 carriers had significantly lower A $\beta$ 42 levels than noncarriers ( $p < 0.05$ ), and a nonsignificant trend level difference in TNF- $\alpha$  ( $p = 0.08$ ). There were no other significant differences between APOE4 carriers and noncarriers as indicated in [Table 1](#).

### 3.2. *TNF- $\alpha$ level is negatively correlated with resting-state functional connectivity*

Linear regression was used to assess the association between higher TNF- $\alpha$  levels and lower FC strength among seed regions and all pairwise ROIs in all subjects ([Fig. 1](#)). To further explore the nature of this relationship, FDR-corrected  $p$ -values were plotted on matrices, split by APOE4 carrier status, revealing a possible E4 effect on the negative correlation between TNF- $\alpha$  and FC ([Fig. 2](#)). Age, sex, education, and diagnosis were included as covariates. The NAcc was identified as the key hub region, defined as the node with the strongest link between all ROIs and TNF- $\alpha$ , as seen in [Fig. 1](#). Specifically, the connectivity between bilateral NAcc and the other 31 individual cortical regions was the best predictor of TNF- $\alpha$  levels across participants. Highly connected regions to NAcc included bilateral posterior middle temporal gyrus (MTG), anterior and posterior superior temporal gyrus, superior temporal gyrus, and temporal pole.

### 3.3. *Moderation effect of APOE4 between TNF- $\alpha$ and FC*

Connectivity between NAcc and posterior and anterior MTG had the strongest relationship with TNF- $\alpha$  ( $p < 0.001$ ). APOE4 was a significant moderator of the relationship between lower FC and higher TNF- $\alpha$  levels. [Fig. 2](#) helps visualize the moderating effect that APOE4 carriers has as seen by the FDR-corrected  $p$ -values along the NAcc column. Specifically, the presence of APOE4 allele moderated the relationship between TNF- $\alpha$  levels and FC in NAcc and posterior MTG (APOE4 carriers: TNF- $\alpha$   $\beta = -1.94$ ,  $t(51) = -4.88$ ,  $p = 0.00001$ ) and NAcc and anterior MTG (APOE4 carriers, TNF- $\alpha$   $\beta = -1.96$ ,  $t(51) = -4.12$ ,  $p = 0.0001$ ) ([Fig. 3](#)).

### 3.4. *Graph theory metrics at the ROI level show differences between APOE4 carriers and noncarriers*

Graph theory metrics for clustering coefficient, average path length, global efficiency, and local efficiency were calculated with the left and right NAcc to characterize the broader ROI connectivity



**Table 1**  
Demographics for the study participants

| Demographics                        | ALL-median (IQR) | APOE4 carriers (n = 34) |                 | APOE4 noncarriers (n = 29) |                 |
|-------------------------------------|------------------|-------------------------|-----------------|----------------------------|-----------------|
|                                     |                  | AD                      | CN              | AD                         | CN              |
| Age (y) <sup>b</sup>                | 70.57 (12.93)    | 75.23 (7.97)            | 68.05 (16.59)   | 76.27 (11.50)              | 64.88 (18.08)   |
| Sex (m/f) <sup>a,b</sup>            | 37/26            | 11/7                    | 9/7             | 10/5                       | 7/7             |
| Education (y)                       | 16 (5.5)         | 16 (5)                  | 14 (2.5)        | 14 (5.25)                  | 16 (2)          |
| Diagnosis (AD/control) <sup>a</sup> | 33/30            | 18                      | 16              | 15                         | 14              |
| MMSE <sup>b</sup>                   | 28 (5)           | 25.5 (4.5)              | 29 (2)          | 27 (7.5)                   | 30 (1)          |
| CDR score (0/0.5/1) <sup>a</sup>    | 30/25/8          | 1/13/4                  | 15/1/0          | 0/11/4                     | 14/0/0          |
| Aβ-42 pg/mL <sup>a,c,e</sup>        | 529.667 (312.94) | 319.45 (235.74)         | 581.07 (278.07) | 505.19 (265.59)            | 643.08 (139.18) |
| T-Tau pg/mL <sup>a,b,d</sup>        | 361.9 (412.51)   | 553.18 (373.03)         | 309.51 (292.43) | 468.25 (349.98)            | 202.99 (91.26)  |
| P-Tau pg/mL                         | 61.892 (43.87)   | 78.18 (69.38)           | 60.45 (34.68)   | 72.56 (46.29)              | 49.13 (24.74)   |
| TNF-α pg/mL <sup>e</sup>            | 0.0939 (0.0809)  | 0.107 (0.081)           | 0.11 (0.114)    | 0.09 (0.0518)              | 0.0806 (0.0275) |

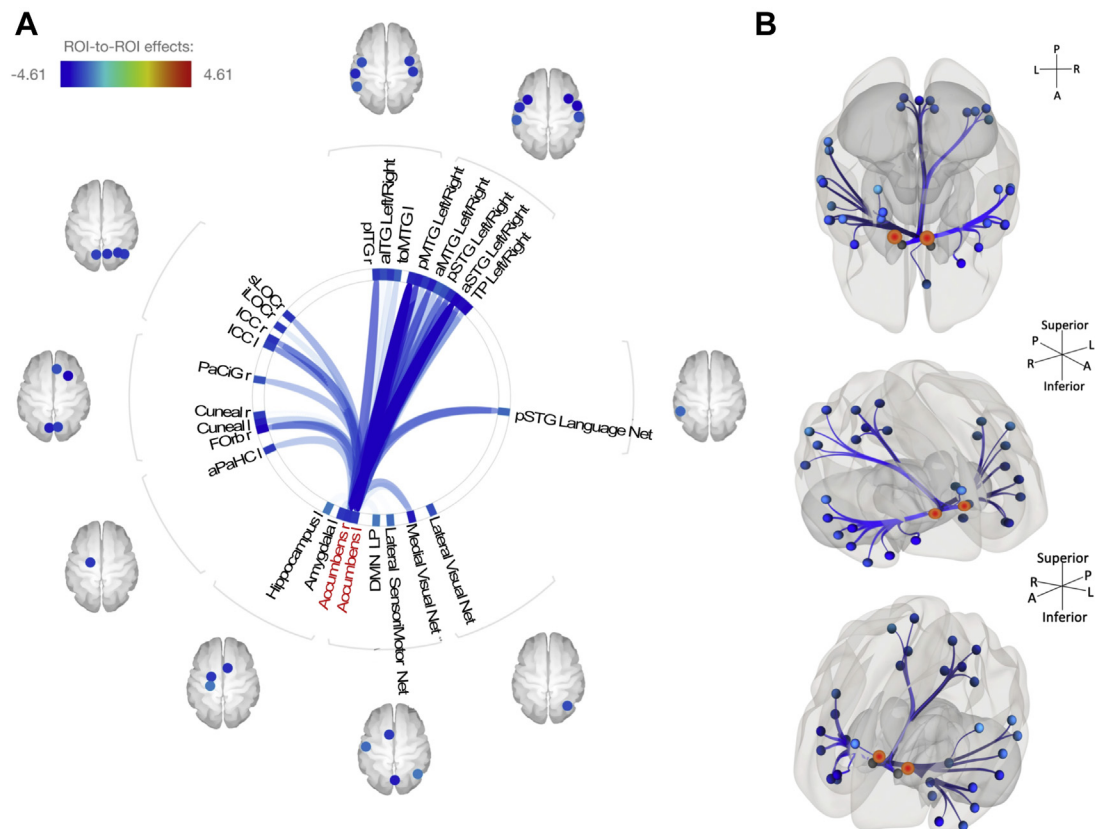
Key: A $\beta$ , amyloid- $\beta$ ; AD, Alzheimer's disease; APOE4, apolipoprotein E- $\epsilon$ 4; CN, cognitively normal; TNF- $\alpha$ , tumor necrosis factor-alpha.

<sup>b</sup> indicates significant difference between AD/CN groups.

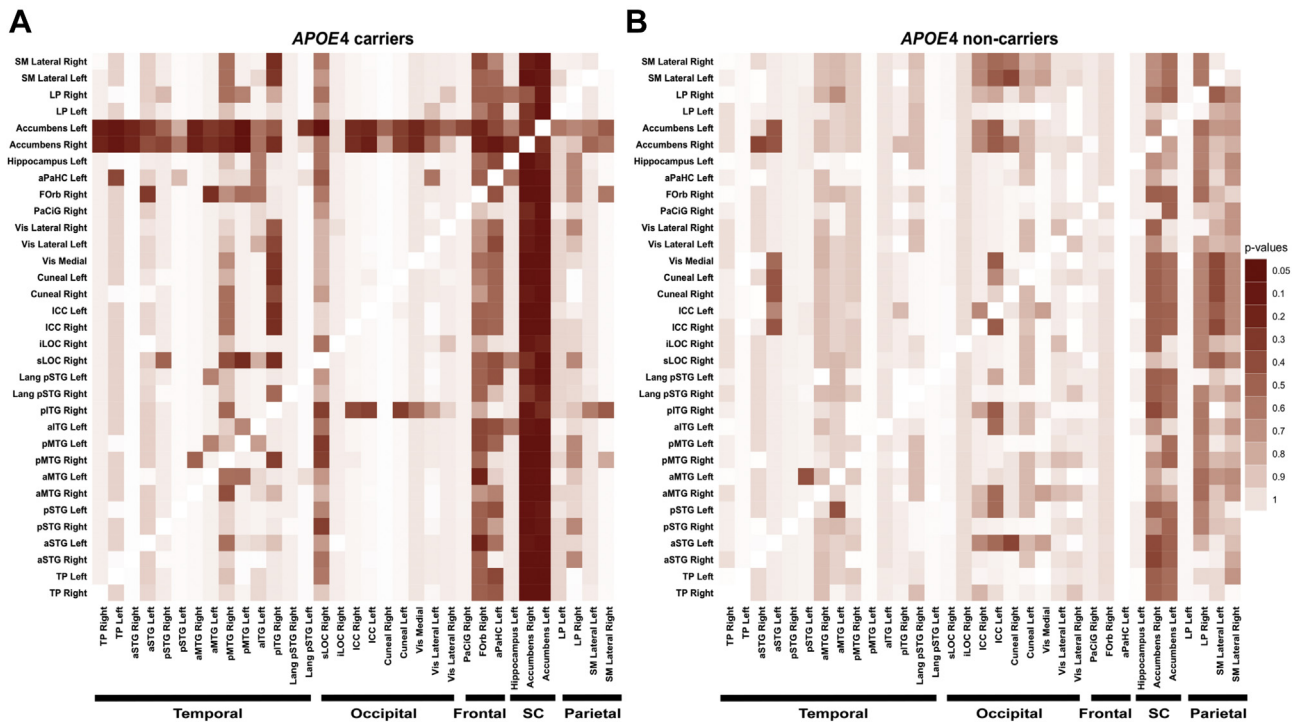
<sup>d</sup> indicates significant differences between APOE4 carrier versus non-carrier subgroups of CN group.

\* indicates significant difference between CN in APOE4 carrier group and AD in APOE4 noncarrier group. APOE4+ participants carried a genotype of APOE3/4. APOE4- participants carried a genotype of APOE3/3.

Clustering coefficient, average path length, global efficiency, and local efficiency were calculated on a whole-brain network level for each participant. A significant difference was found



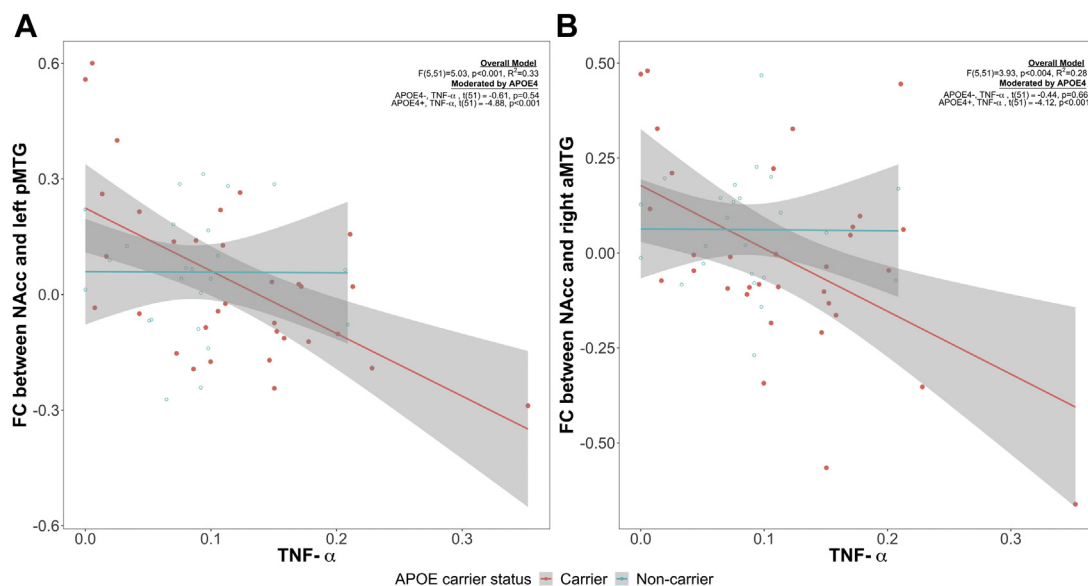
**Fig. 1.** Association between TNF- $\alpha$  and FC. Higher TNF- $\alpha$  levels were correlated with lower FC between NAcc (important for decision-making and inhibitory control) and regions involved in memory, visual processing, and language. (A) Connectogram with blue lines representing significant negative correlations between seed and targets after FDR correction. Opacity and width of line are proportional to FDR-corrected  $p$ -value. (B) Results are visualized anatomically on a brain. Red spheres indicate the NAcc as seed region and blue regions indicate significant ROIs (saturation of blue spheres proportional to  $p$ -value). Abbreviations: TNF- $\alpha$ , tumor necrosis factor-alpha; FC, functional connectivity; NAcc, nucleus accumbens.



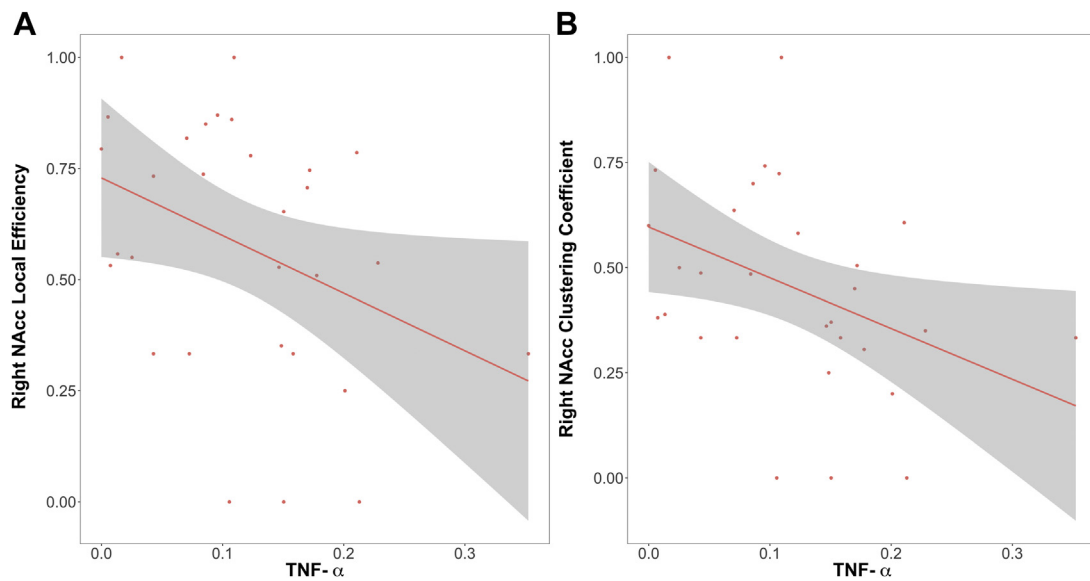
**Fig. 2.** Two-sided  $p$ -value matrices separated by  $APOE4$  carrier status. Matrices, split by  $APOE4$  carrier status, show FDR-corrected  $p$ -values of linear regression model to show the relationship between  $TNF-\alpha$  levels and FC values of ROIs. Connectivity between left and right NAcc and 31 individual cortical regions was significantly associated with  $TNF-\alpha$  levels in  $APOE4$  carriers (A) relative to noncarriers (B) as visualized by the darker red color. Age, sex, education, and diagnosis were included as covariates. Columns represent seed regions; rows represent projection regions. Abbreviations:  $APOE4$ , apolipoprotein E- $\epsilon 4$ ;  $TNF-\alpha$ , tumor necrosis factor-alpha; FC, functional connectivity; NAcc, nucleus accumbens.

between CN and AD participants among all graph theory metrics measuring whole-brain network characteristics (Table 2). Specifically, global efficiency was higher, whereas local efficiency, clustering coefficient, and average path length of the whole-brain network among AD individuals was lower compared with CN participants. In addition, for each metric, whole-brain network visualizations were plotted allowing for further observations

regarding spatial change in network organization as well as plotting the distribution of values per metric by diagnosis to better quantify and characterize the change in network organization (Fig. 5). Variance was calculated to examine the spread of our sample, skewness to determine the symmetry of the sample/distribution, and kurtosis to measure the sharpness of the peak of our distribution (Table 2).



**Fig. 3.** Moderation analysis.  $APOE4$  carrier status moderates the relationship between  $TNF-\alpha$  and functional connectivity of the NAcc to (A) left posterior MTG (pMTG) and to (B) anterior MTG (aMTG).  $APOE4$  carriers had a stronger negative association between  $TNF-\alpha$  and FC. Full model estimates are shown from the moderation analysis. Abbreviations:  $APOE4$ , apolipoprotein E- $\epsilon 4$ ;  $TNF-\alpha$ , tumor necrosis factor-alpha; NAcc, nucleus accumbens.



**Fig. 4.** Connectivity characteristics of NAcc node correlated with TNF- $\alpha$  only in APOE4 carriers. (A) Local efficiency and (B) clustering coefficient in the right NAcc were negatively correlated with TNF- $\alpha$  levels. Specifically, the right NAcc showed less integration with its neighboring ROIs and less clustering when TNF- $\alpha$  levels were high. Global efficiency and average path were not correlated with TNF- $\alpha$ . Abbreviations: APOE4, apolipoprotein E- $\epsilon$ 4; TNF- $\alpha$ , tumor necrosis factor- $\alpha$ ; NAcc, nucleus accumbens.

Within the AD group, overall variance was smaller indicating a loss of network organization as you reduce the inherent hierarchy of nodes (i.e., those acting as hub regions) that is normally present in a healthy brain and consisted of flatter distributions as indicated by a smaller kurtosis value (except in the case of global efficiency). Skewness tended to be more positive in the CN sample, signifying a greater network organizational structure. The difference in path lengths between AD and CN was the most visible (Fig. 5D). The CN group had lengthier paths ( $>2.72$ ) and fewer shorter paths ( $<2.23$ ) compared with the AD group. These shorter path lengths among the AD indicate a less integrated and more modular network pointing to a loss of organization within the AD participants' network.

Overall, local efficiency and clustering coefficient is disrupted in APOE4 carriers relative to noncarriers at the ROI level, whereas all metrics at the global network level were disrupted in AD relative to CN.

#### 4. Discussion

Our study sought to investigate the relationship between TNF- $\alpha$ , an inflammatory CSF cytokine relevant to AD, and rsfMRI FC. In

doing so, we identified a functional connectivity network negatively affected by higher TNF- $\alpha$  levels especially among brain regions known to be important in decision-making and inhibitory control, but with implications for memory, language, and visual processing. On further analysis, this relationship with TNF- $\alpha$  was moderated by APOE4 carrier status. Together, these findings support a heightened proinflammatory environment in APOE4 carriers that impacts functional connectivity of cognitively meaningful networks that include brain regions NAcc and MTG.

The NAcc is central in the functional dopaminergic circuit involving the hippocampus and midbrain and plays a pivotal role in the formation of spatial memory, formation of novelty and persistent memory traces, as well as promoting reward directed behavior (Lisman and Grace, 2005; Roy et al., 2016; Russo and Nestler, 2013). Damage in these areas is presumably the principal cause for the onset of memory loss and both cognitive and noncognitive hallmarks of clinical AD (Squire et al., 2004; Wang and Morris, 2010). In fact, one study implicated that the circuit involving the hippocampus, midbrain and NAcc was severely impaired in an AD mouse model. This impairment resulted from a degeneration of dopaminergic neurons which is essential for correct functioning of the excitatory synaptic transmission between the NAcc and other regions (Cordella et al., 2018). In further support of the importance of the NAcc, another study found that atrophy in the NAcc (via reduced volume) correlated with clinical scores such as the MMSE and MOCA (Nie et al., 2017). Although the studies aforementioned show an important role of the NAcc in AD, our observations showing the relevance of NAcc FC in the context of neuroinflammation add new insight to its role in AD.

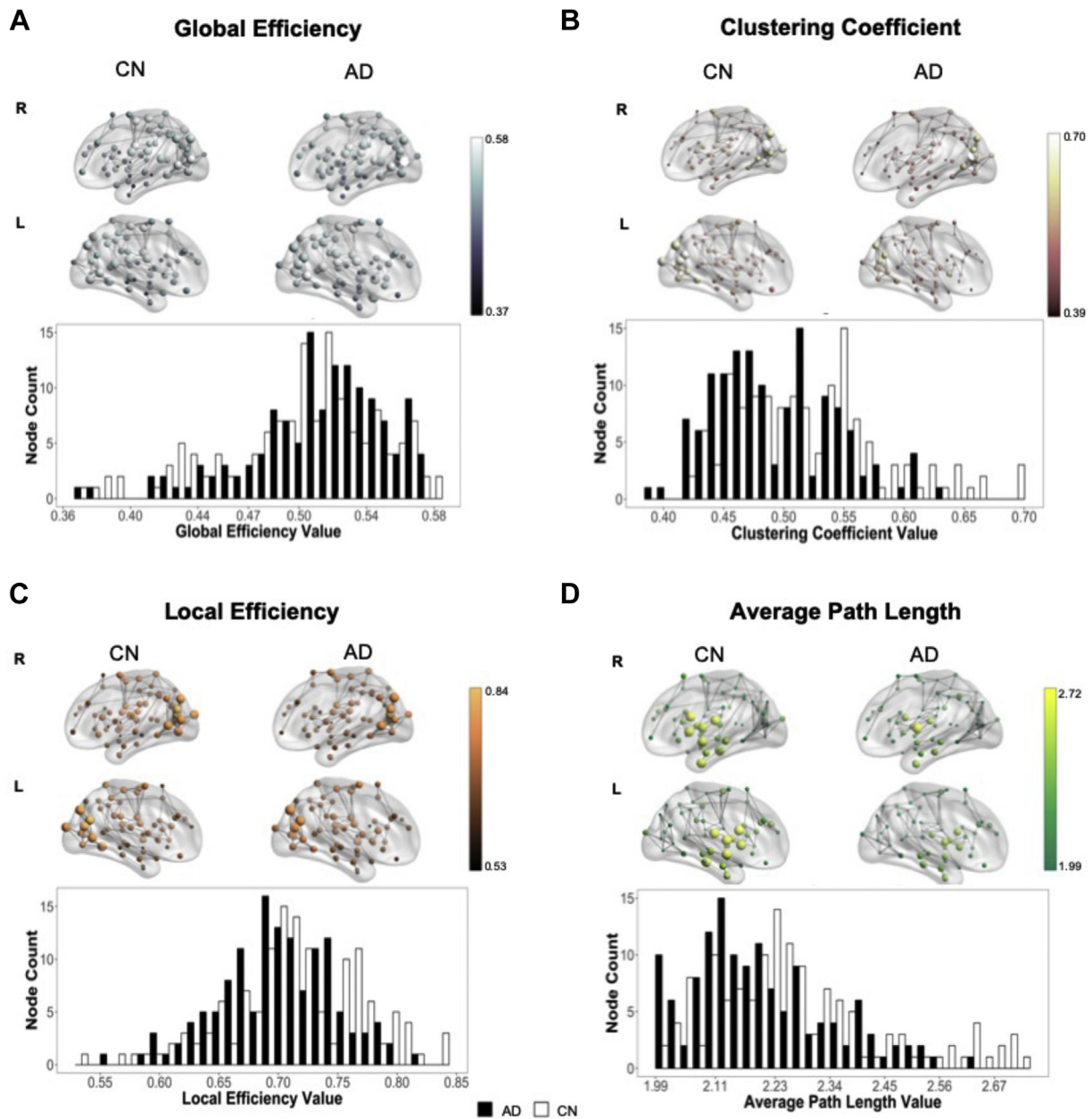
MTG has been implicated across many studies to serve a dynamic role in cognition, including semantic memory processing and language (Cabeza and Nyberg, 2000), observation of motion (Rizzolatti et al., 1996), deductive reasoning (Goel et al., 1998), dynamic facial expressions (Sato et al., 2012), and multimodal and higher sensory processing (Mesulam, 1998). Indeed, it has been proposed that MTG facilitates integration of information between ventral portions of inferior frontal gyrus and regions in the default mode network and executive network (Davey et al., 2016). Therefore, the MTG is spatially well positioned to serve as a hub

**Table 2**  
Connectivity characteristics between CN and AD participants

| Measure                | Group | Mean          | Variance      | Skewness | Kurtosis |
|------------------------|-------|---------------|---------------|----------|----------|
| Global efficiency      | CN    | <b>0.5009</b> | 0.0022        | −0.698   | 3.0281   |
|                        | AD    | <b>0.5125</b> | 0.0017        | −0.9601  | 4.0758   |
| Local efficiency       | CN    | <b>0.7139</b> | <b>0.0033</b> | −0.3135  | 3.1841   |
|                        | AD    | <b>0.7</b>    | <b>0.0023</b> | −0.2049  | 2.9144   |
| Clustering coefficient | CN    | <b>0.5214</b> | <b>0.0038</b> | 0.7414   | 3.1858   |
|                        | AD    | <b>0.4937</b> | <b>0.0025</b> | 0.4945   | 2.7032   |
| Average path length    | CN    | <b>2.2716</b> | <b>0.0303</b> | 0.8843   | 3.2139   |
|                        | AD    | <b>2.2031</b> | <b>0.0195</b> | 0.7247   | 3.0054   |

Bold indicates significance ( $p < 0.05$ ). There was a significant difference in all graph theory measures between AD and CN. There was no difference between APOE4 carriers and noncarriers. We averaged metric ROI values across the brain for each subject to obtain a mean value per diagnosis. Additionally, we used variance, skewness and kurtosis to better characterize the difference in distributions between AD and CN for our respective graph theory metrics.

Key: AD, Alzheimer's disease; APOE4, apolipoprotein E- $\epsilon$ 4; CN, cognitively normal.



**Fig. 5.** Whole-brain network visualizations and distributions for each metric. Each graph theory metric is represented as a whole brain network between AD participants and CN participants. ROIs represent spherical nodes where size and color saturation indicate the numerical value of the metric (i.e., lighter color and larger node represent a larger graph theory metric). (A) Global efficiency values in R and L hemispheres between CN and AD participants. Histogram displays the distribution of the metric by diagnosis. The y-axis indicates how many nodes fall within a certain metric value. Here we see the difference in distributions between CN and AD. (B) Clustering coefficient, (C) local efficiency, and (D) average path. Within the AD group overall variance was smaller indicating a smaller spread of values and consisted of flatter distributions as indicated by a smaller kurtosis value (except in the case of global efficiency). Skewness tended to be greater in the CN sample, indicating a greater network organizational structure (with more high and low values, indicating the importance of certain nodes as compared with others). CN group had lengthier paths and fewer shorter paths compared with the AD group indicating a less integrated and more modular network for AD group. These differences in distribution indicate a loss of organization within the AD participants' network as graph theory metrics show less variance and narrower distribution reflecting a loss of high and low frequencies. Abbreviations: AD, Alzheimer's disease; CN, cognitively normal.

for the default mode network, language, and executive function networks. In support of this, a separate study demonstrated the left MTG as a hub using the participation coefficient (a measure of the distribution of a node's edges among the communities of a network and used to determine a node's "hubness" (Guimerà and Nunes Amaral, 2005)). This study reported that left MTG mediates crosstalk between different functional modules in resting-state networks of those with social anxiety disorder (Yun et al., 2017). Furthermore, it was presumed that poorer cognitive functioning and presence of clinical symptoms (Kambeitz et al., 2016; van den Heuvel and Sporns, 2013; Warren et al., 2014) can result from

lesions within important hub regions such as left MTG (Pannekoek et al., 2013; Qiu et al., 2014, 2015; Shang et al., 2014). Our results, showing strong reduced functional connectivity between and among these 2 important regions lend support to the more "integrated view" that acknowledges the synergy between different brain regions, shifting focus from a "neuron-centric" view which is restricted to local cell dysfunctions (Canter et al., 2016; Palop and Mucke, 2010).

We also examined the functional characteristics of our seed region to the entire network using all brain ROIs (Appendix). Because the NAcc receives information from a variety of systems that



promote distinct patterns of behavior, this region may play a more prominent role in other areas besides reward behavior. Therefore, we used graph theory metrics computing local, clustering, and global efficiency, as well as average path length for the left and right NAcc and overall brain network. We found that the right NAcc showed reduced local efficiency and clustering in *APOE4* carriers with higher inflammation as measured by  $\text{TNF-}\alpha$ . This suggests that the right NAcc showed less integration with its neighboring ROIs and less clustering when  $\text{TNF-}\alpha$  levels were high. Global efficiency and average path length for NAcc were not correlated with  $\text{TNF-}\alpha$ . The NAcc serves as a critical node within the brain's reward circuitry; yet, a more integrated role for this seed region proposed by Mogenson (Mogenson et al., 1993) suggests that the NAcc serves as a limbic-motor interface. Thus, the NAcc may be important for critical high-level tasks, including the selection of actions that help facilitate efficient goal-directed behavior (Floresco, 2015). These goals are set by frontal and temporal lobe regions with which the NAcc interfaces (Floresco et al., 1997; Floresco and Phillips, 1999; Ito et al., 2008; Mannella et al., 2013). Our results contribute to a growing body of literature that has reported differences between *APOE4* carriers and noncarriers using graph theoretical measures at the ROI level (Filippini et al., 2009; Goryawala et al., 2014; Wink et al., 2018), and importantly highlight the relationship between inflammation, functional connectivity, and *APOE4*.

When we compared these graph theory metrics across the whole brain between clinical diagnoses (CN vs. AD), we found significant differences in all measurements, suggesting whole-brain network breakdown in those with clinical AD relative to CN participants. Important to note, the topological organization of a network is directly related to its local and global efficiency (Bullmore and Sporns, 2009; Rubinov and Sporns, 2010), as both are used to determine the brain network's capability of integrating information effectively. For instance, local efficiency provides an indication of how effectively information is integrated between immediate neighbors of a given node, whereas global efficiency provides an indication of how effectively information is integrated across the entire network. We found that while the functional network of participants with AD had a mean global efficiency value that was higher than our CN group among all ROIs, the number of ROIs with a high global efficiency value was less than that of the CN participants. This would indicate that there are fewer ROIs acting as central hub regions in the AD group (Buckner et al., 2009; He et al., 2008; Sporns et al., 2007). Participants with AD also had a greater number of ROIs with low local efficiency values as well as an overall lower local efficiency mean. This difference between global and local efficiency values between AD and CN groups indicates network organizational changes that may result in inadequate or inefficient transfer of information between ROIs, which is consistent with "small-world" organization that reflects the optimal balance between local processing and global integration in the human brain (Bassett and Bullmore, 2016).

Last, taking into consideration both clustering coefficient and path length among ROIs, the network of participants with AD was less integrated and contained a more modular network resulting in reduced path lengths and lower clustering coefficient values. Previous research has shown both resting-state network and modularity network differences between CN and AD groups and at various stages along the disease (Contreras et al., 2019; Sanz-Arigita et al., 2010; Supekar et al., 2008; Tijms et al., 2013; Wang et al., 2010, 2013). These results, taken together, may indicate a possible staging of early network breakdown at the local network level that is related to inflammation and that whole brain network breakdown follows along the spectrum of clinical disease.

Consistent with our findings, other studies have demonstrated higher inflammation and decreased functional connectivity in

other patient populations. For instance, patients with major depressive disorder had reliably elevated cytokine levels, including CRP, interleukins 6, 1 $\beta$ , and 1 that were associated with decreased connectivity within striatal reward-related brain regions (i.e., putamen, caudate, and subdivisions on the striatum) (Felger et al., 2016). Similarly, nonhuman primate work has demonstrated that chronic administration of inflammatory cytokines decreased striatal dopamine release (Felger et al., 2007; Felger and Miller, 2012). Therefore, inflammation-related reductions in corticostriatal connectivity, may serve as a new pathophysiological pathway to explore in an effort to better understand how inflammation effects brain function and connectivity. Another study measured neuroinflammation via activated microglia using a PET radioactive tracer and found that AD-related neuroinflammation in white matter tissue of both mice and humans was increased compared with control mice (Raj et al., 2017). Because white matter tracts in the brain can be related to functional connectivity and speed of processing, compromised white matter integrity during aging can lead to symptomatology such as compromised cognitive performance, as shown by de Lange et al (de Lange et al., 2016). They found differences in white matter integrity correlated to the participants' ability to adapt to demands of cognitive tasks and overall memory performance. In addition to the association between functional brain differences and inflammation, structural MRI studies have demonstrated a relationship between MRI markers of brain volume and CSF and plasma inflammation biomarkers in AD. A study with 42 participants with AD found a correlation between the CSF/serum IgG index and the indices of AD severity, including clinical dementia and medial temporal lobe atrophy (Matsumoto et al., 2008). Another study identified that 34 MRI volumetric brain regions (including the hippocampus, cingulate cortex, amygdala, and basal ganglia) combined with plasma cytokines and chemokines provided the best predictor of conversion from cognitive impairment to AD (Furney et al., 2011). Last, higher levels of systemic inflammation have been associated with reduced fractional anisotropy in white matter and lower cognition (Arfanakis et al., 2013). In a sample of nondemented older participants, systemic inflammation, assessed using a composite measure of inflammatory markers CRP and  $\text{TNF-}\alpha$ , was found to negatively correlate with diffusion anisotropy in the corpus callosum and visual cognition suggesting inflammation is associated with lower microstructural integrity which leads to impaired higher-order visual cognition in aging adults. With evidence mounting on the effects of inflammation on large scale brain changes, it has become even more important to understand the role of inflammation on AD pathophysiology.

The amyloid cascade hypothesis states that deposition of  $\text{A}\beta$  is the causative agent of AD and that neurofibrillary tangles, inflammation, cell loss, vascular damage, and cognitive impairment are downstream effects of  $\text{A}\beta$ , with interactions between events (Hardy and Higgins, 1992; Selkoe, 1991). Yet, mounting evidence suggests that neuroinflammation may play a substantial role in Alzheimer's pathogenesis. While the interaction and temporal sequence of senile plaques, neurofibrillary tangles, and neuroinflammation remain inconclusive, animal models suggest neuroinflammation may contribute to the pathogenesis of plaques and tangles themselves (Zhang et al., 2013). This is underlined by findings that genes for immune receptors, such as *TREM2*, a microglial activation marker (Guerreiro et al., 2013) and *CD33*, also a microglial activation marker and thought to inhibit  $\text{A}\beta$  clearance (Bradshaw et al., 2013; Griciu et al., 2013), are associated with AD. Analysis of clinical manifestations, which precede AD, such as mild cognitive impairment, further argues for substantial involvement of inflammation in the pathogenesis of AD.

#### 4.1. Limitations and future directions

There are limitations to the present study. For instance, we were limited in that our *APOE4* carriers were all heterogenous E3/4 carriers whereas our noncarriers were homozygous E3/3 carriers. Ideally, we would have liked to be able to parse out effects for those that were homozygous for E4 and any protective effects that E2 homozygous and heterozygous might have. It's also worth noting that we elected to use a specific brain parcellation scheme that also identifies ROIs belonging to particular resting state networks. It is possible that alternative anatomic and functional templates could return different results. Thus, future studies including multiple brain parcellation schemes and resting state network definitions would help determine the validity of our findings. We also acknowledge that better spatial resolution and improved acquisition parameters can greatly improve data quality as well as improve the stability and reproducibility of FC ROI-to-ROI results. Although this was not possible within this study, owing to the retrospective nature of the data, future studies will use superior methodology. In addition, we did not examine the role of other prominent inflammatory markers that may be modulating TNF- $\alpha$  such as IL-8, IL-6, as well as determining if blood-brain barrier (BBB) breakdown influences these relationships. We also recognize we were limited by the availability of clinical assessments as is the case with retrospective data. As much as 52% of clinical data were either missing, not collected, or incomplete. However, we are currently collecting additional data with similar variables (i.e., resting state fMRI data with advanced acquisition sequences, *APOE* genotype information availability, cognitive data) as well as CSF data that include a wide range of inflammatory markers and markers to measure BBB breakdown. These data will enable future studies to explore other markers and better establish a biological pathway for TNF- $\alpha$  (i.e., establish if vascular damage, via measuring BBB breakdown, leads to an increase in inflammatory markers).

In summary, we have provided evidence implicating neuroinflammation in *APOE4* and AD, showing that higher CSF TNF- $\alpha$  levels are associated with lower FC between brain regions important in decision-making and inhibitory control in *APOE4* carriers. Investigating inflammation and functional connectivity in older adults at risk for AD may help us better understand how the immune system contributes to the pathogenesis of AD.

#### Disclosure

The authors have no conflict of interest to disclose.

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#### Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.neurobiolaging.2019.10.013>.

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